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Amendments to the Claims

1. (currently amended) A method for the treatment of a disorder of the eye comprising: administering to a subject a therapeutically effective amount of a composition comprising a dsRNA between 21 and 23 nucleotides in length and a carrier, said dsRNA having a nucleotide sequence corresponding to mRNA of SEQ ID NO: 3 a target gene expressed in the eye; said administering of the composition occurring outside the blood-retina barrier, and said composition inhibiting the target gene by RNA interference inside the eye.

2.-3. (canceled)

- 4. (previously presented) The method of claim 1, wherein said disorder is related to angiogenesis and/or neovascularization.
- 5. (previously presented) The method of claim 1, wherein said disorder is related to the retinal pigment epithelium (RPE), neurosensory retina, choroid, and a combination thereof.
- 6. (previously presented) The method of claim 1, wherein said disorder is wet age-related macular degeneration (AMD) or diabetic retinopathy.
- 7.-8. (canceled)
- 9. (currently amended) The method of claim 1, wherein said dsRNA is an inhibitor of expression of <u>SEQ ID NO: 3</u> the target gene.
- 10. (currently amended) The method of claim 9, wherein said inhibitor inhibits expression of the target gene SEQ ID NO: 3 involved in angiogenesis and/or neovascularization.
- 11.-15. (canceled)
- 16. (previously presented) The method of claim 1, wherein the dsRNA comprises a terminal 3'-hydroxyl group.
- 17.-93. (canceled)
- 94. (withdrawn) The method of claim 1, further comprising preparing the dsRNA.

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95: (withdrawn) The method of claim 1, further comprising diagnosing a subject with a disorder or a predisposition to a disorder of the eye.

- 96. (currently amended) The method of claim 1, further comprising detecting a product of SEQ ID NO: 3 the target gene.
- 97. (withdrawn) The method of claim 1, further comprising isolating the target gene.
- 98. (previously presented) The method of claim 1, wherein said administering is by systemic administration.